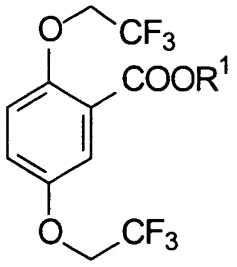
IN THE CLAIMS

Please amend the claims as follows:

1. (previously presented) Process for the preparation of Flecainide, as Flecainide base or any pharmaceutically acceptable salts thereof, comprising:

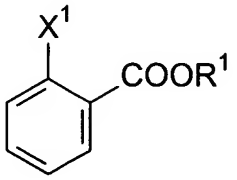
preparation of a compound of formula VI



(VI)

wherein R^1 is H, alkali metal or a C_1 to C_9 alkyl group;

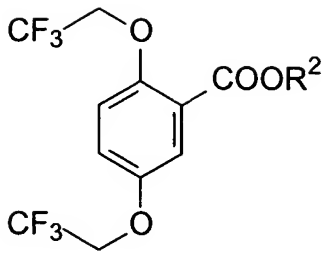
from compounds of formula III



(III)

wherein X^1 is F, Cl, Br or I;

optional conversion of the compound of formula VI to the ester of formula VII by reacting with a hydroxyl compound R^2OH;



(VII)

wherein R^2 is C_1 to C_9 alkyl group, aryl group or succinimidyl;

amide formation of the compound of formula VI or VII forming flecainide base by reacting with 2-(aminomethyl)piperidine and;

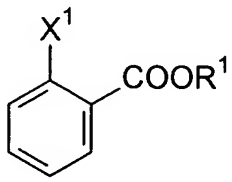
optionally forming a pharmaceutically acceptable salt thereof.

2. (original) The process of Claim 1 wherein the amide formation is selective.

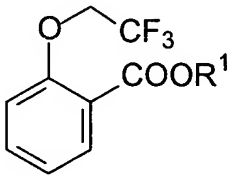
3. (cancelled)

4. (previously presented) Process for the preparation of Flecainide, as Flecainide base or any pharmaceutically acceptable salts thereof, comprising

reaction of the 2-halobenzoic acid derivatives of formula III with an alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethanol in the presence of a copper-containing catalyst in a solvent to form 2-(2,2,2-trifluoroethoxy)benzoic acid derivatives of formula IV;



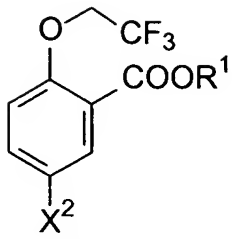
(III)



(IV)

wherein X^1 is F, Cl, Br or I and R^1 is H, alkali metal or a C_1 to C_9 alkyl group;

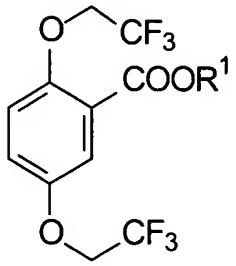
halogenation of the compounds of formula IV to form 5-halo-2-(2,2,2-trifluoroethoxy)benzoic acid derivatives of formula V;



(V)

wherein X^2 is Cl, Br, or I.

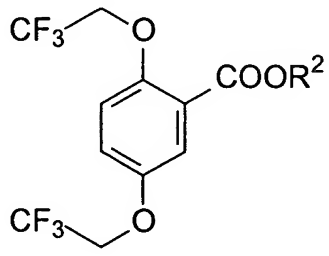
reaction of the compounds of formula V with an alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethanol in the presence of a copper-containing catalyst in a solvent to form compounds of formula VI;



(VI)

wherein R^1 is H, alkali metal or a C_1 to C_9 alkyl group;

optional conversion of the compounds of formula VI to a new ester of formula VII by reacting with hydroxyl compound R^2OH;



(VII)

wherein R^2 is C_1 to C_9 alkyl group, aryl group or succinimidyl;

selective amide formation by reacting compounds of formula VI or VII with 2-(aminomethyl)piperidine forming flecainide base;

optionally forming a pharmaceutically acceptable salt thereof.

5. (original) The process of Claim 4 wherein either solvent comprises a polar solvent.

6. (original) The process of Claim 4 wherein the pharmaceutically acceptable salt is the monoacetate salt.

7. (original) The process according to Claim 4, wherein the alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethoxide is sodium, potassium, calcium or lithium 2,2,2-trifluoroethoxide.

8. (previously presented) The process according to Claim 4, wherein the alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethanol is synthesized by reacting 2,2,2-trifluoroethanol with a base selected from potassium *tert*-butoxide, sodium *tert*-butoxide, sodium isopropoxide and sodium methoxide.

9. (cancelled)

10. (previously presented) The process according to Claim 4 wherein the copper-containing catalyst comprises cupric chloride, cupric bromide, cupric iodide, cuprous chloride, cuprous bromide, cuprous iodide, copper (I) oxide, copper (II) oxide or copper-zinc alloy.

11. (original) The process according to Claim 4, wherein X^2 is Br.

12. (original) The process according to Claim 4, wherein R^2 is selected from methyl, ethyl, benzyl and phenyl.

13. (original) The process according to Claim 4, wherein the compound of formula VI or VII is 2,5-bis-(2,2,2-trifluoroethoxy)benzoate.

14.(original) The process according to Claim 13 wherein any of the reactions is carried out in aliphatic, cycloaliphatic or aromatic solvents from 5 to 10 carbon atoms or ethers from 4 to 10 carbon atoms.

15. (previously presented) The process according to Claim 14, wherein the solvents comprise hexane, heptane, cyclohexane, tetrahydrofuran, 1,2-domethoxyethane, diethyleneglycol dimethyl ether, toluene, xylene, or acetonitrile.

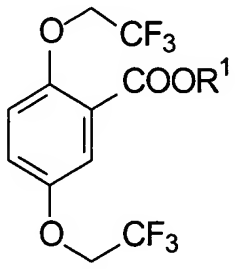
16. (original) The process according to Claim 13, wherein the reaction temperature is between 0°C to 150°C.

17. (original) The process according to Claim 13, wherein the temperature is between 50°C to 120°C.

18. (original) The process according to Claim 13, wherein the molar ratio between 2,5-bis-(2,2,2-trifluoroethoxy)benzoate and 2-aminomethylpiperidine is from 1:1 to 1:2.

19. (original) The process according to Claim 18, wherein the molar ratio is from 1:1 to1:1.5.

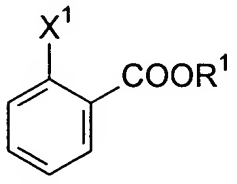
20 (previously presented) The process for the preparation benzoic acid derivatives of formula VI ;



(VI)

wherein R^1 is H, alkali metal or a C_1 to C_9 alkyl group;

from compounds of formula III

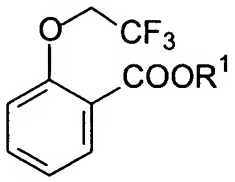


(III)

wherein X^1 is F, Cl, Br or I;

comprising:

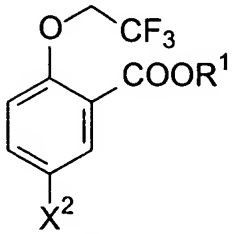
reaction of compounds of formula III with an alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethanol in the presence of a copper-containing catalyst in a solvent to form compounds of formula IV;



(IV)

wherein R^1 is H, alkali metal or a C_1 to C_9 alkyl group;

halogenation of the compounds of formula IV to form compounds of formula V;



(V)

wherein X^2 is Cl, Br, or I[[.]];

reaction of compounds of formula V with an alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethanol in the presence of a copper-containing catalyst in a solvent.

21. (original) The process according to Claim 20 wherein either solvent comprises a polar solvent.

22. (original) The process according to Claim 20, wherein the alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethoxide is sodium, potassium, calcium or lithium 2,2,2-trifluoroethoxide.

23. (previously presented) The process according to Claim 20, wherein the alkali or alkaline earth metal alkoxide of 2,2,2-trifluoroethoxide is synthesized by reacting 2,2,2-triflouroethanol with a base

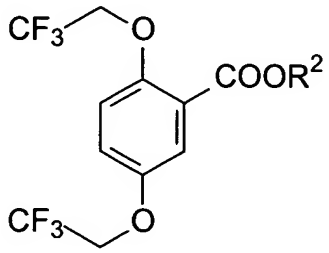
selected from potassium *tert*-butoxide, sodium *tert*-butoxide, sodium isopropoxide and ~~or~~ sodium methoxide.

24. (cancelled)

25. (previously presented) The process according to Claim 20 wherein the copper-containing catalyst comprises cupric chloride, cupric bromide, cupric iodide, cuprous chloride, cuprous bromide, cuprous iodide, copper (I) oxide, copper (II) oxide, or copper-zinc alloy.

26. (original) The process according to Claim 20, wherein X^2 is Br.

27. (currently amended) The process for the preparation of Flecainide from 2,5-bis(2,2,2-trifluoroethoxy)benzoic acid derivatives of formula VII,



(VII)

wherein R^2 is ~~C_1 to C_9 alkyl group, aryl group or succinimidyl and wherein R^2 is not 2,2,2-trifluoroethyl or cynomethyl groups~~ methyl, ethyl, propyl, butyl, benzyl, phenyl or succinimidyl;

comprising the selective amide formation by reacting the benzoic acid derivative of formula VII with 2-(aminomethyl)piperidine.

28. (original) The process according to Claim 27, wherein the reaction is carried out in aliphatic, cycloaliphatic or aromatic solvents from 5 to 10 carbon atoms or ethers from 4 to 10 carbon atoms.

29.(currently amended) The process according to Claim ~~28~~ 27, wherein the reaction is carried out in solvents and the solvents are selected from hexane, heptane, cyclohexane, tetrahydrofuran, 1,2-demethoxyethane, diethyleneglycol dimethyl ether, toluene, xylene, acetonitrile.

30. (previously presented) The process according to Claim 28, wherein the solvent is toluene or xylene.

31. (original) The process according to Claim 27, wherein the reaction temperature is between 0°C and 150°C.

32. (original) The process according to Claim 27, wherein temperature range is between 50°C and 120°C.

33. (original) The process according to Claim 27, wherein the molar ratio between the benzoic acid derivative and 2-aminomethylpiperidine is from 1:1 to 1:2.

34. (original).The process according to Claim 33, wherein the molar ratio is from 1:1 to 1:1.5.

35. (original) 5-Bromo-2-(2,2,2-trifluoroethoxy)benzoic acid.